



06/01/22 Morning Report with @CPSolvers



Case Presenter: Steph (@StephVSherman) Case Discussants: Zaven (@sargsyanz) and the chat <3

<p>CC: 2 week leg ulcer</p> <p>HPI: 28yo W - 3 years ago p/w with chest pain and rash. She was told she might have SLE. Never been on therapy. Over the last 2 years, R>L leg swelling on and off. When the swell occurs, she has groin lymphadenopathy. When the swelling goes off, so does the lymphadenopathy. Over the last month, R leg pretty swollen. Last 2 weeks with first episode of painful ulcer in the chin. Came to the ED for evaluation.</p>	<p>Vitals: T: nl HR: nl BP: nl RR: nl SpO₂: nl</p> <p>Exam:</p> <p>Gen: well-appearing</p> <p>HEENT: nl CV: nl Pulm: nl Abd: nl</p> <p>Extremities/Skin: R>L pitting edema ("puffy")</p> <p>R shin - 4 cm well-circumscribed tender necrotic ulcer with black eschar</p>	<p>Problem Representation: 28yoW with a PMH significant for suspicious SLE p/w 2 weeks R leg ulcer and edema. Found to have tender necrotic ulcer with black eschar and labs consistent w/ IDA and elevated ESR and CRP. Plan film of the leg revealed diffuse sclerotic lesions.</p>								
<table border="1"> <tr> <td data-bbox="21 633 157 722">PMH: none</td> <td data-bbox="157 633 436 722">Fam Hx: Non contributory</td> </tr> <tr> <td data-bbox="21 722 157 852">Meds: none</td> <td data-bbox="157 722 436 852">Soc Hx: 4yo healthy child Works in an office</td> </tr> <tr> <td colspan="2" data-bbox="21 852 436 966">Health-Related Behaviors: Non contributory</td> </tr> <tr> <td colspan="2" data-bbox="21 966 436 1177">Allergies: None</td> </tr> </table>	PMH: none	Fam Hx: Non contributory	Meds: none	Soc Hx: 4yo healthy child Works in an office	Health-Related Behaviors: Non contributory		Allergies: None		<p>Notable Labs & Imaging:</p> <p>Hematology:</p> <p>WBC: nl Hgb: 7.5 (low MCV 62) Plt: nl</p> <p>Iron deficiency anemia</p> <p>Chemistry:</p> <p>BMP - unremarkable HIV neg</p> <p>Urinalysis - 30 mg/dl protein, no RBCs.</p> <p>PTT: 39 (corrected with mixing study) Normal INR</p> <p>Leg US - no clots</p> <p>ESR >100 CRP 13</p> <p>Cr nl BUN nl</p> <p>Heavy menstrual bleeding Positive ANA</p> <p>Biopsy - reactive lymphadenopathy with normal cytometry</p> <p>2 transient face and lips swelling episodes - 4 years ago - never happened again</p> <p>Hb electrophoresis - nl</p> <p>CT venography - no compression / diffuse bony sclerotic lesions</p> <p>Plan film leg - diffuse sclerosis lesions</p> <p>CXR nl</p>	<p>Teaching Points (Seyma & Kirtan):</p> <ul style="list-style-type: none"> ● Leg Ulcers: Vascular (Art/Venous/Vasculitis), Infection, Malignancy, Autoimmune (PAN, Lupus, Sarcoid, SVV, APLAS or pyoderma gangrenosum) <ul style="list-style-type: none"> ● Varicose veins predispose to vascular ulcers w/ unilateral edema ● Unilateral leg edema: venous insufficiency, lymph edema (e.g. elephantiasis due to filariasis), lymphangitis, cellulitis, DVT, ruptured baker's cyst, pelvic neoplasm ;usually not a/w LAD (reactive?) <ul style="list-style-type: none"> ● May Thurner Anomaly/Cockett syndrome: L Iliac vein compressed by R iliac artery ● Black eschar: <ul style="list-style-type: none"> ● Infection: rickettsia, scrub typhus, anthrax, tularemia, fungal; calciphylaxis (uremic calcific arteriopathy) or ecthyma gangrenosum (sepsis w/ pseudomonas aeruginosa), gas gangrene ● Vasculitides: <ul style="list-style-type: none"> ● Medium vessel: nodules, ulcers; Small-vessel: petechiae, palpable purpura ● Neutrophilic dermatoses: Sweet, Pyoderma g. → initially rather erythematous, then ulcerate ● Microcytic anemia: iron-def (menstruation, etc), Hb-pathies (sickle-cell disease), <ul style="list-style-type: none"> ● Mentzer-Index: RBC/MCV-Index (>13 → rather iron deficiency mediated anemia; <13 β-thalassemia) ● "TAILS" Mnemonic for microcytic anemia: Thalassemia, Anemia of chronic dz, Iron deficiency, Lead poisoning, Sideroblastic ● >50% pyoderma gangrenosum: idiopathic; otherwise a/w autoimmune, sarcoid ● Isolated elevated PTT: think of APLS ● Sclerotic bone lesions: malignancy (multiple myeloma, metastases: prostate-cancer, infiltrative (Langerhans cell histiocytosis, Gaucher's dz, Erdheim Chester) Autoinflammatory dz (Schnitzler), Mastocytosis, POEMS, metabolic (Paget's dz) ● >90% of patients w/ sarcoid should have an abnormal Chest CT scan (LAD, ...)
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